MA

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SKILLS

- Neurology trials
- Patient assessment
- Data analysis
- Regulatory compliance
- Team collaboration
- Clinical research

EDUCATION

MASTER OF SCIENCE IN NEUROSCIENCE, UNIVERSITY OF BRAIN SCIENCES, 2013

LANGUAGE

- English
- Spanish
- German

ACHIEVEMENTS

- Contributed to a trial that led to the identification of a new treatment for Alzheimer's disease.
- Received the 'Research Excellence' award for outstanding contributions to neurology trials.
- Increased patient enrollment rates by 50% through community outreach initiatives.

Michael Anderson

CLINICAL TRIALS SCIENTIST

Detail-oriented Clinical Trials Scientist specializing in neurology with over 9 years of experience in clinical trial design and execution. Proven expertise in managing trials for neurodegenerative diseases, including Alzheimer's and Parkinson's. Strong background in data analysis and patient interaction, ensuring ethical standards are met throughout the trial process. Excellent organizational skills and the ability to work collaboratively with interdisciplinary teams.

EXPERIENCE

CLINICAL TRIALS SCIENTIST

NeuroResearch Institute

2016 - Present

- Managed multi-center phase II trials for Alzheimer's treatment, overseeing patient recruitment and data collection.
- Conducted regular monitoring visits to ensure compliance with GCP and FDA regulations.
- Coordinated with neurologists and research staff to facilitate seamless trial operations.
- Utilized statistical software for data analysis, presenting findings to the research team.
- Engaged with patients to assess treatment effects and gather qualitative feedback.
- Maintained comprehensive trial documentation to support regulatory submissions.

CLINICAL RESEARCH ASSISTANT

Alzheimer's Research Foundation

2014 - 2016

- Supported the management of phase I trials for Alzheimer's drugs, assisting with patient assessments and data entry.
- Maintained regulatory documentation in compliance with institutional guidelines.
- Conducted site visits to monitor patient safety and data accuracy.
- Assisted in the development of study protocols and informed consent documents.
- Engaged with participants to provide information and support throughout the trial process.
- Contributed to the preparation of regulatory submissions, facilitating timely approvals.